# IN THE UNITED STATES PATENT AND TRADEMARK OFFICE BOARD OF PATENT APPEALS AND INTERFERENCES

In re Application of: Robert E. Arbogast et al. Confirm. No.: 4457

Application No.: 09/893,535 <u>Examiner</u>: Dilek B.

Cobanoglu

OHI 1717-008A

Attorney

Docket No.:

Filing Date: 06/29/2001 Art Unit: 3626

Title: System, Method, And

Computer Program Product

For Configuring And Purchasing A Medical

Device

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/Trisha Beachy-Bryant/ Trisha Beachy-Bryant (paralegal)

# APPELLANTS BRIEF UNDER 37 CFR § 41.37

#### Dear Sir:

This is an appeal from a final rejection mailed on August 08, 2006, in which the Examiner rejected all of the claims pending in the present application. The requisite \$250.00 (small entity) filing fee set forth in 37 CFR § 41.20(b)(2) accompanies this Brief.

**REAL PARTY IN INTEREST** 

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Inventor: Arbogast et al.

Examiner: Dilek B. Cobanoglu

Ford, Phillip Lee Harrison, Raymond Francis, Keith W. Justus, Rebecca L. Halley,

Bradley A. Spitzer, Thomas D. Chamberlain and Eric L. Kershner are currently named

as inventors on the present application. The present application has been assigned to

the Ohio Willow Wood Company, which is the real party of interest in the present appeal.

RELATED APPEALS AND INTERFERENCES

There are no other prior or pending appeals, interferences or judicial proceedings

that relate to or would in any way affect or have a bearing on the Board's decision in the

present appeal.

STATUS OF CLAIMS

Claims 1-39, 46-49, 65-69 and 80-82 are pending in the present application.

Claims 1-39, 46-49, 65-69 and 80-82 stand rejected. Claims 40-45, 50-64, 70-79 and

83-85 have been withdrawn from consideration as the result of a previously issued

restriction requirement. Claims 1-39, 46-49, 65-69 and 80-82 are the subject of this

appeal.

STATUS OF AMENDMENTS

Per the Advisory Action of December 18, 2006, Appellant's November 08, 2006

response to the Examiner's final Office Action mailed on August 08, 2006 was entered

for purposes of appeal, along with an explanation as to how the amended claims would

be rejected.

## SUMMARY OF CLAIMED SUBJECT MATTER

The claimed subject matter is directed to an automated system and method of configuring a medical device, such as a prosthesis, from a collection of individual medical device components. More specifically, based on one or a few criteria (typically patient attributes and/or desires), a system and method of the present invention can query one or more databases containing relevant components, retrieve and sort through the multitudes of components available with respect to a particular medical device (e.g., a prosthetic leg or arm) and provide one or a number of acceptable prosthesis configurations using various combinations of qualifying components. Such criteria may include, for example, patient weight, patient activity level, medical device cost, or medical device weight. Thus, a system and method of the present invention can greatly reduce the time required to configure a medical device — which would currently be accomplished by a practitioner using a number of different catalogues, etc.

Support for the various elements of independent claims 1, 31, 46 and 65, as presented in Appellant's November 08, 2006 response to the Examiner's final Office Action, is identified in the charts below:

	Claim 1	
Element Number	Claim Elements from Claim 1	Support in Application
1	A system for configuring a medical device, comprising:	At least paragraphs [0001], [0016]- [0023], [0039]-[0108]; FIGS. 1-2, 7-8, 10-11.
2	a digital repository populated with entries defining a plurality of medical device components,	At least paragraphs [0041], [0049]-[0050], [0061], [0065] and [0078]; Abstract.

3	each entry associated with an individual medical device component and having	At least paragraphs [0049]-[0050], [0061] and [0065].
4	a component identification indicator,	At least paragraphs [0041], [0049]-[0050], [0061], [0065], [0072] and [0078].
5	a component class indicator, and	At least paragraphs [0065], [0063] and [0078]; FIG. 3.
6	at least one patient attribute indicator;	At least paragraphs [0018]-[0019], [0024], [0057], [0062] and [0083].
7	a processor; and	At least paragraphs [0097], [0100]- [0101], and [0105]-[0106]; FIG. 11.
8	a computer readable medium encoded with processor readable instructions that when executed by the processor implement	At least paragraphs [0098], [0101]- [0103] and [0105]; FIG. 11.
9	a practitioner user interface mechanism configured to provide a practitioner with access to entries in the digital repository via a network and to allow the practitioner to provide at least one patient interview answer indicator,	At least paragraphs [0021], [0039], [0048], [0053]-[0054], [0058]-[0062], [0070], [0077]-[0078], [0089]; FIGS. 1, 9 and 10A-10H.
10	a patient interview mechanism configured to receive over the network the at least one patient interview answer indicator corresponding to an attribute of a patient and to store the at least one patient interview answer indicator in a memory, and	At least paragraphs [0051]-[0052], [0058], [0061]-[0062]; FIGS. 4, 6 and 9.
11	a configurator mechanism configured to select a subset of entries from the digital repository based on the at least one patient interview answer indicator in the memory,	At least paragraphs [0054], [0058], [0068]-[0069], [0071]-[0072]; FIGS. 2, 4, 4A 6 and 9.

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the subset of entries including entries corresponding to individual medical device components that collectively form a medical device meeting a need of the patient.	At least paragraphs [0005], [0018], [0020], [0024], [0050], [0068]- [0069], [0071]-[0072]; FIGS. 2, 4, 4A 6 and 9.
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	Claim 31	
Element Number	Claim Elements from Claim 31	Support in Application
1	A method for configuring a medical device, comprising the steps of:	At least paragraphs [0001], [0016]- [0023], [0039]-[0108]; FIGS. 3-10.
2	populating a digital repository with information corresponding to a plurality of medical device components;	At least paragraphs [0041], [0049]-[0050], [0061], [0065] and [0078]; Abstract.
3	interviewing a patient having a need for a medical device to determine at least one patient attribute;	At least paragraphs [0021], [0051]- [0052], [0061][0062], [0067], [0081], [0090]; FIG. 3.
4	storing the at least one patient attribute in a memory; and	At least paragraphs [0018]-[0019], [0024], [0057], [0062] and [0083].
5	querying the digital repository for a subset of medical device components based on the at least one patient attribute,	At least paragraphs [0054], [0058], [0068]-[0069], [0071]-[0072]; FIGS. 2, 4, 4A, 6 and 9.
6	the subset of medical device components collectively forming a medical device meeting the need of the patient.	At least paragraphs [0005], [0018], [0020], [0024], [0050], [0068]- [0069], [0071]-[0072]; FIGS. 2, 4, 4A, 6 and 9.

	Claim 46	
Element Number	Claim Elements from Claim 46	Support in Application
1	A system for configuring a medical device, comprising:	At least paragraphs [0001], [0016]-[0023], [0039]-[0108]; FIGS. 1-2, 7-8, 10-11.
2	means for populating a digital repository with information corresponding to a plurality of individual medical device components;	At least paragraphs [0041], [0049]- [0050], [0061], [0065] and [0078]; Abstract.
3	means for interviewing a patient having a need for a medical device to determine at least one patient attribute;	At least paragraphs [0021], [0051]- [0052]; FIGS. 1 and 10A.
4	means for storing the at least one patient attribute in a memory; and	At least paragraphs [0018]-[0019], [0024], [0057], [0062], [0083], [0102]-[0103], [0105]-[0106].
5	means for querying the digital repository for a subset of medical device components based on the at least one patient attribute,	At least paragraphs [0054], [0058], [0068]-[0069], [0071]-[0072]; FIGS. 2, 4, 4A, 6 and 9.
6	the subset of medical device components collectively forming a medical device meeting the need of the patient.	At least paragraphs [0005], [0018], [0020], [0024], [0050], [0068]- [0069], [0071]-[0072]; FIGS. 2, 4, 4A, 6 and 9.

Claim 65		
Element Number	Claim Elements from Claim 65	Support in Application
1	A method for configuring a medical device, comprising the steps of:	At least paragraphs [0001], [0016]- [0023], [0039]-[0108]; FIGS. 3-10.

2	populating a digital repository with information corresponding to a plurality of individual medical device components;	At least paragraphs [0041], [0049]-[0050], [0061], [0065] and [0078]; Abstract.
3	populating the digital repository with patient historical information associated with a patient;	At least paragraphs [0023], [0050], [0072]; Abstract.
4	interviewing the patient having a need for a medical device to determine at least one patient attribute;	At least paragraphs [0021], [0051]- [0052], [0061][0062], [0067], [0081], [0090]; FIG. 3.
5	storing the at least one patient attribute in a memory via a digital communication link;	At least paragraphs [0018]-[0019], [0024], [0043], [0045]-[0046], [0051], [0057], [0062], [0083], [0107]-[0108]; FIGS. 1 and 9.
6	querying the digital repository for a subset of medical device components based on the at least one patient attribute,	At least paragraphs [0054], [0058], [0068]-[0069], [0071]-[0072]; FIGS. 2, 4, 4A, 6 and 9.
7	the subset of medical device components collectively forming a medical device meeting the need of the patient;	At least paragraphs [0005], [0018], [0020], [0024], [0050], [0068]- [0069], [0071]-[0072]; FIGS. 2, 4, 4A, 6 and 9.
8	ordering the medical device over the digital communication link; and	At least paragraphs [0017], [0019], [0024]-[0025], [0029], [0031]-[0032], [0034], [0041], [0062], [0067], [0070], [0076]-[0078], [0085]-[0088]; FIGS. 1, 3-4, 5-6, 10G-10H.
9	storing information corresponding to the medical device in the digital repository associated with the patient.	At least paragraphs [0077], [0085], [0088].

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**GROUNDS OF REJECTION TO BE REVIEWED ON APPEAL** 

With respect to Claim 1:

(1) Whether Clynch (US 6,463,351) in view of DeBusk et al. (US 6,581,204)

teaches or suggests a system for *configuring* a medical device;

(2)whether Clynch in view of DeBusk et al. teaches or suggests a digital

repository (database) of medical device *components*:

(3)whether Clynch in view of DeBusk et al. teaches or suggests a component

class indicator or at least one patient attribute indicator:

(4) whether Clynch in view of DeBusk et al. teaches or suggests a patient

interview mechanism or allows a practitioner to supply at least one patient

interview answer indicator, or stores such a patient interview answer indicator

in memory: and

whether Clynch in view of DeBusk et al. teaches or suggests configuring a (5)

medical device from a collection of medical device components, the

components selected from a plurality of possible qualifying components based

on some defined patient need(s).

With respect to Claim 31:

(1) Whether Clynch teaches a method for *configuring* a medical device;

(2)whether Clynch teaches populating a digital repository (database) with

information relating to a multitude of medical device components;

(3)whether Clynch teaches interviewing a patient, or interviewing a patient to

obtain information later used in configuring the medical device;

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(4) whether Clynch subsequently teaches storing information obtained from the

patient interview;

(5) whether Clynch teaches searching the database for data on a plurality of

individual medical device components, the search criteria based on patient

information obtained in the patient interview; and

(6) whether Clynch teaches configuring a medical device from a collection of

medical device components, the components selected from a plurality of

possible qualifying components based on some defined patient need(s).

With respect to Claim 46:

(1) Whether Clynch in view of DeBusk et al. teaches or suggests a system for

configuring a medical device;

(2) whether Clynch in view of DeBusk et al. teaches or suggests a digital

repository (database) of medical device components and a means of placing

data in such a repository;

(3) whether Clynch in view of DeBusk et al. teaches or suggests interviewing a

patient in general or interviewing a patient to obtain information later used in

configuring the medical device and, if so, if a means for effecting such an

interview is also taught;

(4) whether Clynch in view of DeBusk et al. subsequently teaches or suggests a

means for storing information obtained from the patient interview;

(5) whether Clynch in view of DeBusk et al. teaches or suggests a means for

searching the database for data on a plurality of individual medical device

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components, the search criteria based on patient information obtained in the

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patient *interview*; and

(6)whether Clynch in view of DeBusk et al. teaches or suggests configuring a

medical device from a collection of medical device components, the

components selected from a plurality of possible qualifying components based

on some defined patient need(s).

With respect to Claim 65:

(1) Whether Clynch teaches a method for *configuring* a medical device:

(2)whether Clynch teaches populating a digital repository (database) with

information relating to a multitude of *individual* medical device *components*;

(3)whether Clynch teaches saving historical information relating to a patient in

the database with the medical device data:

(4) whether Clynch teaches interviewing a patient, or interviewing a patient to

obtain information later used in configuring the medical device;

(5)whether Clynch subsequently teaches storing information obtained from the

patient interview:

(6)whether Clynch teaches searching the database for data on a plurality of

individual medical device components, the search criteria based on patient

information obtained in the patient interview; and

(7)whether Clynch teaches configuring a medical device from a collection of

medical device components, the components selected from a plurality of

possible qualifying components based on some defined patient need(s);

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(8) whether Clynch in any way teaches ordering a medical device, and ordering a

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medical device over a digital communication link; and

(9) whether Clynch teaches storing information about the configured medical

device in the database with the patient data.

**ARGUMENT** 

Rejection of Claims 31-37, 39, 46-48, 65-67 and 82 Under 35 U.S.C. § 102(e)

The Examiner rejected claims 31-37, 39, 46-48, 65-67 and 82 under 35 U.S.C. §

102(e) as being anticipated by Clynch (US 6,463,351). Appellants respectfully disagree

with the Examiner's rejection of said claims.

Clynch does not teach the subject matter of the rejected claims. Rather, Clynch

is directed specifically and only to producing an individual custom prosthetic socket or

other similar interface component. Specifically, Clynch teaches an improved system

and method for producing a customized interface (e.g., socket) for a prosthetic or

orthotic device. The specific steps required to produce such a socket were set forth in

Appellants' November 08, 2006 response to the Examiner's final Office action and can

be found in Clynch at col. 2, II. 28-48; col. 2, line 58 - col. 3, line 26; claim 1; and Figure

Therefore, Clynch teaches only the manufacturing of an individual prosthetic

component, not the configuring, producing or assembling of a complex medical device

(e.g., prosthesis or orthosis) from a *collection* of *individual* medical device *components*.

While Clynch does teach a method for manufacturing a prosthetic socket, it fails

to teach a new method for designing, configuring or assembling the remainder of a

prosthesis (of which the Clynch interface is merely a part). Rather, the remainder of the

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prosthesis or orthosis must still be configured by a known manual method like the

present invention is specifically designed to render obsolete. That is, once a prosthetist.

orthotist or other practitioner uses the system and method of Clynch to obtain a

customized prosthetic or orthotic socket, the prosthetist/orthotist must still manually

search for the remainder of the components required to assemble the overall prosthesis

or orthosis from a variety of catalogs and other sources and then manually sort through

a potentially enormous number of possible combinations of said components to arrive at

one or several acceptable configurations. It is this type of manual process that the

method of the rejected claims renders obsolete.

That is, based on one or a few criteria, a system and method of the present

invention can sort through one or more databases containing the multitudes of

components available with respect to a particular prosthesis type (e.g., leg, arm) or

other medical device type and provide one or a number of acceptable prosthesis or

other medical device configurations using various combinations of qualifying

components. Such criteria may include, for example, a prosthesis of the lowest cost or

of the lightest weight, and/or may be based on patient-specific data such as weight,

activity level, etc.

In this regard. Appellants further assert that Clynch also fails to teach the

interviewing of a patient. Clynch teaches only that a practitioner interacts with a patient

during the casting and scanning of the patient's residual limb. There is no teaching that

the practitioner interviews the patient, formally or otherwise, to obtain at least one

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patient attribute that will be subsequently used in the configuration and selection of a

prosthesis. Rather, the Examiner merely infers that this is likely. Appellants disagree.

Throughout the prosecution of the present application, the Examiner has

continued to assert that particular sections of Clynch teach the claimed aspects of the

present invention. Appellants have, on several occasions, made known to the Examiner

that they have been unable to locate support for the Examiner's assertions at the cited

locations of Clynch. These concerns have not, however, been adequately addressed

by the Examiner. For example, the following discrepancies were pointed out in

Appellants' November 08, 2006 response to the Examiner's final Office action:

With respect to independent claim 31 of the present application, the Examiner

asserts that the claimed subject matter

querying the digital repository for a subset of medical device

components based on the at least one patient attribute, the subset of

medical device components collectively forming a medical device

meeting the need of the patient

is taught by Clynch at col. 7, II, 22-44 and 61-65, as well as col. 4, II, 14-39 thereof.

However, Clynch at column 4 merely describes that prosthetists, orthopedic surgeons,

podiatrists, radiologists and plastic surgeons may practice the invention in creating

prosthetic devices, orthotic devices, etc. This is no doubt true - but the invention of

Clynch is the creation of a custom medical device interface, not an overall prosthesis.

Thus, in order to configure the remainder of the medical device, these practitioners are

currently forced to employ the manual system that the present invention renders

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obsolete. Further, Appellants have been unable to locate any reference to a subset of

medical device components, a digital repository, or the querying of a digital repository

anywhere in the cited portion of column 4.

Similarly, there is absolutely no mention of a subset of medical device

components, a digital repository of medical device components, or querying a digital

repository for a subset of medical device components in column 7, II. 22-44 or 61-65 of

Clynch. By referring to line 14 of column 7, it is clear that the language cited by the

Examiner describes only the digital image manipulation portion of the Clynch invention.

(See also Fig. 3, referred to therein). Everything described in these sections of Clynch,

and shown in Fig. 3, is related to a practitioner manipulating the digital image of the cast

body part to produce areas of relief or build-up in the subsequently manufactured

interface component. (See col. 1, II. 33-38 for more general description). The

referenced shrink and smooth operations, as well as the database storage of default

modifications (i.e., pre-defined image changes), are all functions of image manipulation.

Even the language of lines 47-52 refers to modifications that may be made to the digital

image in order to produce specific modifications to the finished interface component in

the stated areas - not to medical device components. There is no discussion of a digital

repository of medical device components, because Clynch is not concerned with

anything more than a single component at a time. Further, the database referred to by

the Examiner stores only information on previous changes made to similar digital body

part (model) images, not information on medical devices.

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With respect to independent claim 46 of the present application, the Examiner

makes the same assertions discussed above with respect to claim 31, and further

asserts that the claimed subject matter

means for populating a digital repository with information corresponding

to a plurality of individual medical device components

is taught by Clynch at col. 4, II. 14-39 and 49-53; and col. 7, II. 61-63 thereof. The lack

of any such disclosure at col. 4, II. 14-39 and col. 7, II. 61-63 of Clynch has already been

addressed above. Col. 4, II. 49-53 of Clynch is similarly lacking in such teachings.

Rather, this section of Clynch discloses only that a scan facility may be equipped with

equipment that allows for the capturing (via scanning) of a 3-dimensional image of a

target (body part) surface, and for conversion of the scanned data into a 3-dimensional

image (see Fig. 3) that may be displayed on a computer. Once again, the cited sections

of Clynch appear to be wholly devoid of any reference to a digital repository or to

medical device components.

The Examiner makes the same assertions with respect to independent claim 65 as

were made with respect to claim 46. As the deficiencies of these cited sections have

already been discussed above, there is no need to repeat the arguments here.

As can be understood from the foregoing, Clynch fails to teach at least several

elements of each of the rejected claims. Consequently, Appellants once again

respectfully submit that Clynch cannot support a rejection of claims 31-37, 39, 46-48,

65-67 and 82 under 35 U.S.C. § 102(e).

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Rejection of Claims 1-5, 8-14, 16, 19, 20, 22-30 and 80-81 35 U.S.C. § 103(a)

The Examiner rejected claims 1-5, 8-14, 16, 19, 20, 22-30 and 80-81 under 35

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U.S.C. § 103(a) as being unpatentable over Clynch in view of DeBusk et al. (US

6,581,204). Appellants respectfully disagree with the Examiner's rejection of said

claims.

Most of the deficiencies of Clynch combined with DeBusk et al. have already

been addressed above. DeBusk et al. appears to be nothing more than an advanced

medical supply inventory tracking and management system and, as such, does nothing

to make up for the above-documented deficiencies of Clynch. In fact, the Examiner

appears to have cited DeBusk et al. only for its asserted disclosure of a component

identification indicator. Therefore, with respect to independent claim 1, Clynch in view

of DeBusk et al. fails to teach or suggest at least:

(1) a digital repository populated with entries defining a plurality of medical device

components, the entries each associated with an *individual* medical device component:

(2) at least one patient attribute indicator associated with the entries;

(3) a practitioner user interface mechanism configured to provide a practitioner

with access to entries in the digital repository via a network and to allow the practitioner

to provide at least one patient interview answer indicator;

(4) a patient interview mechanism configured to receive over the network the at

least one patient interview answer indicator corresponding to an attribute of a patient

and to store the at least one patient interview answer indicator in a memory; and

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(5) a configurator mechanism configured to select a subset of entries from the

digital repository based on the at least one patient interview answer indicator in the

memory, the subset of entries including entries corresponding to individual medical

device components that collectively form a medical device meeting a need of the

patient.

Furthermore, as discussed with respect to the previous rejection, the Examiner

has continued to cite sections of Clynch in which Appellants have been unable to find

various allegedly present subject matter. For example, the following specific

discrepancies were pointed out in Appellants' November 08, 2006 response to the

Examiner's final Office action with respect to claim 1:

With respect to independent claim 1 of the present application, the Examiner asserts

that the claimed subject matter

a digital repository populated with entries defining a plurality of medical

device components, each entry associated with an individual medical

device component

is taught by Clynch at col. 4, Il. 14-39 and lines 49-53 as well as col. 7, line 61 to col. 8,

line 10 thereof, and that the claimed subject matter

a configurator mechanism configured to select a subset of entries from

the digital repository based on the at least one patient interview answer

indicator in the memory, the subset of entries including entries

corresponding to individual medical device components that collectively

form a medical device meeting a need of the patient

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is taught by Clynch at col. 7, Il. 22-44 and 61-65. The teachings of Clynch at column 4,

II. 14-39 and lines 49-53 have been previously addressed. Clynch at col. 7, line 61 to

col. 8, line 10, merely expounds slightly on the digital image manipulation discussion

described above. The digital image manipulation of Clynch is in no way related to the

sorting and selection of multiple medical device components contemplated by the

present invention. The cited sections of Clynch, like the remainder of said reference.

are simply devoid of such teaching or suggestion, and the Examiner does not appear to

have cited Debusk in the rejection of claim 1.

As with the previous rejection, Clynch in view of DeBusk et al. fails to teach at

least several elements of each of the rejected claims. Therefore, Appellants once again

respectfully submit that Clynch in view of DeBusk et al. cannot support a rejection of

claims 1-5, 8-14, 16, 19, 20, 22-30 and 80-81 under 35 U.S.C. § 103(a).

Other Discrepancies

In addition to the discrepancies already pointed out above with respect to the

cited sections of Clynch, Appellants also disagree with the Examiner's reliance on the

following sections of Clynch and assert that the language thereof does not support the

Examiner's conclusions:

- in paragraph 11, the "body portions" referred to by the Examiner are human

body portions that may be scanned and imaged according to the invention of Clynch.

They are not medical devices nor medical device components and, thus, their relevance

is not apparent to Appellants:

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- in paragraph 12, the patellar tenon bar, head of fibula and posterior wall

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extension referred to by the Examiner as "medical components" and appearing in col. 7.

II. 44-52 of Clynch, are in actuality various anatomical features of an amputated lower

leg. These are human body part features, *not* "medical components";

- in paragraph 13, the tubular modeling material referred to by the Examiner is

not a medical device or a medical device component. Rather, the modeling material is

a casting material that is temporarily placed on a patient's body part to produce the 3-

dimensional model that is eventually scanned, digitized and manipulated. As the

modeling material does not remain on the patient's body, nor become part of the

resulting interface component (socket), there would be no reason for the practitioner or

anyone else to ask the patient whether the tubular modeling material is comfortable or

suits the patient's lifestyle (as seems to be asserted by the Examiner):

- in paragraph 14, the scan data file referred to by the Examiner is not a stored

patient attribute (weight, height, activity level, prosthesis weight preference, etc.).

Rather, the file the Examiner refers to is simply a saved digital image of the cast model,

either before or after manipulation by a practitioner;

- in paragraph 15, Appellants reassert the same argument: operating a CAD

software package to manipulate a 3-dimensional image of a body part (i.e., to produce

areas of build-up or relief in the finished interface component) is clearly not the same as

querying a database of medical device components in relation to selecting acceptable

components for the construction of a prosthesis. Shape/size manipulation of a 3-

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dimensional image is simply not the same as, or even related to, sorting and selectively

choosing components from a database of components based on predetermined criteria:

- in paragraph 16, Appellants again stress that this section of Clynch teaches

only that an interface component created according to the described invention may be

subsequently used in the assembly of, e.g., a prosthesis or orthosis. However, it is not

taught or suggested that the invention can be used to create an overall prosthesis or

orthosis. Rather, the manual configuration technique that the present invention seeks to

render obsolete would currently have to be practiced in order to create a complete

prosthesis or orthosis. Further, the invention of Clynch is directed only to the interface

component of such a device. As the remainder of such a device does not contact the

patient, it should be obvious that the invention of Clynch would not be applicable to

other components; and

- in paragraph 21, the pull down menu options referred to by the Examiner are

nothing more than menus that allow for a practitioner to perform particular

manipulations of the 3-dimensional digital image of the modeled body part (much like

the pull down menus of any CAD software package allow for the specific manipulation

of a drawing model). Contrary to the Examiner's belief, these menus nor the software

associated therewith, have anything whatsoever to do with selecting a subset of medical

device components from a digital repository thereof.

CONCLUSION

For at least the foregoing reasons, it is submitted that the Examiner's rejection of

claims 1-39, 46-49, 65-69 and 80-82 is unsupported by the cited references. As such,

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reversal of the Examiner's rejection of claims 1-39, 46-49, 65-69 and 80-82 and allowance of the present application is respectfully requested.

Respectfully submitted,

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CLAIMS APPENDIX

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(as appearing in Appellants' November 08, 2006 response)

Claim 1 (previously presented): A system for configuring a medical device, comprising:

a digital repository populated with entries defining a plurality of medical device

components, each entry associated with an individual medical device component and

having

a component identification indicator,

a component class indicator, and

at least one patient attribute indicator:

a processor; and

a computer readable medium encoded with processor readable

instructions that when executed by the processor implement

a practitioner user interface mechanism configured to provide a practitioner with

access to entries in the digital repository via a network and to allow the practitioner to

provide at least one patient interview answer indicator,

a patient interview mechanism configured to receive over the network the at least

one patient interview answer indicator corresponding to an attribute of a patient and to

store the at least one patient interview answer indicator in a memory, and

a configurator mechanism configured to select a subset of entries from the digital

repository based on the at least one patient interview answer indicator in the memory.

the subset of entries including entries corresponding to individual medical device

components that collectively form a medical device meeting a need of the patient.

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Claim 2 (original): The system of Claim 1, wherein the medical device comprises at

least one of a lower extremity prosthetic device, an upper extremity prosthetic device, a

lower extremity orthotic device, an upper extremity orthotic device, and a spinal orthotic

device.

Claim 3 (original): The system of Claim 1, wherein:

the computer readable medium is further encoded with processor readable

instructions that when executed by the processor further implement a customization

mechanism configured to at least one of add, remove, and modify at least one entry of

the subset of entries selected by the configurator mechanism, and the practitioner user

interface mechanism is further configured to provide access to the customization

mechanism.

Claim 4 (original): The system of Claim 3, wherein the practitioner user interface

mechanism is further configured to provide a summary page of components customized

by the customization mechanism.

Claim 5 (original): The system of Claim 1, wherein:

the computer readable medium is further encoded with processor readable

instructions that when executed by the processor further implement a medical device

shopping mechanism configured to place an order for the medical device and to store

order information in the digital repository, and

the practitioner user interface mechanism is further configured to provide access

to the medical device shopping mechanism.

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Claim 6 (original): The system of Claim 5, wherein the medical device shopping

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mechanism is further configured to determine all applicable price discounts for the

medical device available to the practitioner.

Claim 7 (original): The system of Claim 1, wherein at least a portion of the practitioner

user interface mechanism is accessible via a personal data assistant.

Claim 8 (original): The system of Claim 1, wherein at least a portion of the network

comprises an Internet protocol based network.

Claim 9 (original): The system of Claim 1, wherein a least a portion of the network is the

Internet.

Claim 10 (original): The system of Claim 1, wherein the digital repository comprises:

a central digital repository, and

a practitioner local digital repository remote from the central database.

Claim 11 (original): The system of Claim 10, wherein:

at least one of the practitioner local digital repository and the central digital

repository is further populated with patient historical entries, the patient historical entries

each associated with an individual patient and having a patient identification indicator,

and at least one patient history indicator.

Claim 12 (original): The system of Claim 11, wherein the at least one patient history

indicator comprises information corresponding to a medical device of the individual

patient.

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Claim 13 (original): The system of Claim 12, wherein the information corresponding to a

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medical device of an individual patient comprises an identification number of a

component of the medical device.

Claim 14 (original): The system of Claim 11, wherein:

the patient historical entries further have at least one patient care indicator.

Claim 15 (original): The system of Claim 14, wherein the patient care indicator

comprises reimbursement information.

Claim 16 (original): The system of Claim 15, wherein the reimbursement information

comprises an L code indicator.

Claim 17 (original): The system of Claim 11, wherein the digital repository is configured

to interface with an external system.

Claim 18 (original): The system of Claim 17, wherein the external system comprises at

least one of a patient management system, a billing system, and an insurance

reimbursement system.

Claim 19 (original): The system of Claim 1, wherein:

the computer readable medium is further encoded with processor readable

instructions that when executed by the processor further implement a patient letter of

necessity generation mechanism configured to generate a letter of necessity for the

patient based on information stored in the digital repository and to store the letter of

necessity in the digital repository, and

the practitioner user interface mechanism is further configured to provide access

to the patient letter of necessity generation mechanism.

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Claim 20 (original): The system of Claim 1, wherein the digital repository comprises a

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database.

Claim 21 (original): The system of Claim 1, wherein the practitioner user interface is

further configured to accept the at least one patient interview answer indicator from an

external device.

Claim 22 (original): The system of Claim 21, wherein the external device is at least one

of a digitizer, a digital camera, and a digital video camera.

Claim 23 (original): The system of Claim 1, wherein:

the entries in the digital repository further have a ranking indicator, and

the configurator mechanism is further configured to select a plurality of subsets of

entries from the digital repository based on the at least one patient interview answer

indicator in the memory, each of the plurality of subsets including entries corresponding

to individual medical device components of a medical device meeting the need of the

patient and being ranked based on the ranking indicator of the entries.

Claim 24 (original): The system of Claim 23, wherein the ranking indicator comprises at

least one of a component cost indicator, a component weight indicator, a component

height indicator, a component width indicator, a component activity level indicator, and

an inventory indicator.

Claim 25 (original): The system of Claim 23, wherein:

the computer readable medium is further encoded with processor readable

instructions that when executed by the processor further implement a customization

mechanism configured to select one of the plurality of subsets of entries and at least

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one of add, remove, and modify at least one entry of the one of the plurality of subsets

of entries selected by the configurator mechanism, and

the practitioner user interface mechanism is further configured to provide access

to the customization mechanism.

Claim 26 (original): The system of Claim 25, wherein:

the computer readable medium is further encoded with processor readable

instructions that when executed by the processor further implement a medical device

shopping mechanism configured to place an order for the medical device corresponding

to the one of the plurality of subsets of entries selected by the customization mechanism

and to store order information in the digital repository, and

the practitioner user interface mechanism is further configured to provide access

to the medical device shopping mechanism.

Claim 27 (original): The system of Claim 25, wherein the practitioner user interface

mechanism is further configured to provide a summary page of components customized

by the customization mechanism.

Claim 28 (previously presented): The system of Claim 1, wherein:

the computer readable medium is further encoded with processor readable

instructions that when executed by the processor further implement a catalog

mechanism configured to select a subset of entries from the digital repository based on

a query and to provide the subset of entries to the practitioner user interface

mechanism, and

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a medical device component shopping mechanism is configured to place an

order for a medical device component corresponding to at least one selected entry of

the subset of entries and to store order information in the digital repository,

the practitioner user interface mechanism further configured to accept the query

from a user, to provide the query to the catalog mechanism, and to select the at least

one selected entry of the subset of entries provided by the catalog mechanism.

Claim 29 (original): The system of Claim 28, wherein:

the computer readable medium is further encoded with processor readable

instructions that when executed by the processor further implement a customization

mechanism configured to select at least one of the subsets of entries and at least one of

add, remove, and modify at least one entry of the one of the plurality of subsets of

entries selected by the catalog mechanism, and

the practitioner user interface mechanism is further configured to provide access

to the customization mechanism.

Claim 30 (original): The system of Claim 29, wherein the practitioner user interface

mechanism is further configured to provide a summary page of components customized

by the customization mechanism.

Claim 31 (previously presented): A method for configuring a medical device, comprising

the steps of:

populating a digital repository with information corresponding to a plurality of

medical device components;

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interviewing a patient having a need for a medical device to determine at least

one patient attribute:

storing the at least one patient attribute in a memory; and

querying the digital repository for a subset of medical device components based

on the at least one patient attribute, the subset of medical device components

collectively forming a medical device meeting the need of the patient.

Claim 32 (original): The method of Claim 31, wherein the medical device comprises at

least one of a lower extremity prosthetic device, an upper extremity prosthetic device, a

lower extremity orthotic device, an upper extremity orthotic device, and a spinal orthotic

device.

Claim 33 (original): The method of Claim 31, further comprising the step of:

customizing at least one of the subset of medical device components to create a

customized medical device further meeting the need of the patient.

Claim 34 (original): The method of Claim 31, wherein the guerving step comprises:

querying the digital repository for a plurality of subsets of medical device

components based on the at least one patient attribute, the subset of medical device

components corresponding to a medical device meeting the need of the patient; and

ranking the plurality of subsets based on a ranking criteria.

Claim 35 (original): The method of Claim 34, wherein the ranking criteria is at least one

of a weight of the medical device, a height of the medical device, a width of the medical

device, a cost of the medical device, an activity level supported by the medical device,

and an inventory status of the medical device.

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Claim 36 (original): The method of Claim 34, further comprising the step of:

selecting one of the plurality of subsets:

customizing the one of the plurality of subsets to create a customized medical

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device further meeting the need of the patient; and

ordering the customized medical device.

Claim 37 (original): The method of Claim 36, wherein the ordering step comprises

reviewing the customized medical device prior to ordering.

Claim 38 (original): The method of Claim 36, wherein the ordering step comprises

determining all applicable price discounts for the medical device for the practitioner.

Claim 39 (original): The method of Claim 31, wherein the interviewing step comprises

entering the at least one patient attribute via at least one of a personal data assistant, a

digitizer, a digital camera, and a digital video camera.

Claim 40 (withdrawn): A method for outsourcing the fabrication of a medical device,

comprising the steps of:

populating a digital repository with information corresponding to a plurality of

medical device components;

interviewing a patient having a need for a medical device to determine at least

one patient attribute:

measuring a body part of the patient and producing a digital map of the body

part; and

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querying the digital repository by a supplier for a subset of medical device

components based on the at least one patient attribute, the subset of medical device

components corresponding to a medical device meeting the need of the patient.

Claim 41 (withdrawn): The method of Claim 40, wherein the medical device comprises

at least one of a lower extremity prosthetic device, an upper extremity prosthetic device,

a lower extremity orthotic device, an upper extremity orthotic device, and a spinal

orthotic device.

Claim 42 (withdrawn): The method of Claim 40, further comprising the step of:

sending the at least one patient attribute and the digital map to the supplier over

a digital communication link.

Claim 43 (withdrawn): The method of Claim 40, further comprising the step of:

customizing the medical device by the supplier based on the digital map to

produce a customized medical device.

Claim 44 (withdrawn): The method of Claim 42, wherein at least a portion of the digital

communication link comprises the Internet.

Claim 45 (withdrawn): The method of Claim 42, further comprising the step of:

shipping the customized medical device to the practitioner.

Claim 46 (previously presented): A system for configuring a medical device, comprising:

means for populating a digital repository with information corresponding to a

plurality of individual medical device components;

means for interviewing a patient having a need for a medical device to determine

at least one patient attribute;

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means for storing the at least one patient attribute in a memory; and

means for querying the digital repository for a subset of medical device

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components based on the at least one patient attribute, the subset of medical device

components collectively forming a medical device meeting the need of the patient.

Claim 47 (original): The system of Claim 46, wherein the medical device comprises at

least one of a lower extremity prosthetic device, an upper extremity prosthetic device, a

lower extremity orthotic device, an upper extremity orthotic device, and a spinal orthotic

device.

Claim 48 (original): The system of Claim 46, further comprising:

means for customizing at least one of the subset of medical device components

to create a customized medical device further meeting the need of the patient.

Claim 49 (original): The system of Claim 46, further comprising:

means for determining applicable discounts for the medical device for the

practitioner.

Claim 50 (withdrawn): A computer program product, comprising:

a computer storage medium and a computer program code mechanism

embedded in the computer storage medium for causing a processor to facilitate the

management of a digital workflow, the computer program code mechanism having

a first computer code device configured to populate a digital repository with data

that is descriptive of components of a medical device,

a second computer code device configured to collect patient information and to

store patient specific information in a memory, and

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a third computer code device configured to select a subset of components stored

in the digital repository based on the patient specific information so as to configure a

medical device meeting a need of a particular patient.

Claim 51 (withdrawn): The computer program product of Claim 50, wherein the medical

device comprises at least one of a lower extremity prosthetic device, an upper extremity

prosthetic device, a lower extremity orthotic device, an upper extremity orthotic device,

and a spinal orthotic device.

Claim 52 (withdrawn): The computer program product of Claim 50, wherein:

the first computer code device is further configured to populate ranking

information associated with the components of a medical device in the digital repository

corresponding to a ranking criteria; and

the third computer code device is further configured to select a plurality of

subsets of components stored in the digital repository based on the patient specific

information and to rank the plurality of subsets based on the ranking criteria so as to

configure a plurality medical devices meeting the need of a particular patient.

Claim 53 (withdrawn): The computer program product of Claim 50, further comprising:

a fourth computer code device configured to customize at least one component

of the subset of components selected by the third computer code device.

Claim 54 (withdrawn): The computer program product of Claim 52, further comprising:

a fourth computer code device configured to customize at least one component of one

of the plurality of subsets of components selected by the third computer code device.

Claim 55 (withdrawn): The computer program product of Claim 50, further comprising:

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a fourth computer code device configured to accept an order for the medical

device over a digital communication link.

Claim 56 (withdrawn): The computer program product of Claim 55, wherein at least a

portion of the digital communication link is the Internet.

Claim 57 (withdrawn): The computer program product of Claim 50, wherein:

the second computer code device is further configured to store the patient

specific information in the memory over a digital communication link.

Claim 58 (withdrawn): The computer program product of Claim 57, wherein at least a

portion of the digital communication link is the Internet.

Claim 59 (withdrawn): The computer program product of Claim 50, wherein:

the second computer code device is further configured to collect patient

information via at least one of a personal data assistant, a personal computer, a

digitizer, a digital camera, and a digital video camera.

Claim 60 (withdrawn): The computer program product of Claim 50, further comprising:

a fourth computer code device configured to provide an interface to an external

system, wherein the first computer code device is further configured to populate patient

historical information associated with a particular patient.

Claim 61 (withdrawn): The computer program product of Claim 59, wherein the patient

historical information comprises an identification number of a component of a medical

device of the patient.

Claim 62 (withdrawn): The computer program product of Claim 60, wherein:

the patient historical information comprises patient care information, and

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the external system comprises at least one of a patient management system, a

billing system, and an insurance reimbursement system.

Claim 63 (withdrawn): The computer program product of Claim 62, wherein the patient

care information comprises at least one of reimbursement information and L code

information.

Claim 64 (withdrawn): The computer program product of Claim 55, wherein the fourth

computer code device is further configured to determine applicable discounts for the

medical device for the practitioner.

Claim 65 (previously presented): A method for configuring a medical device, comprising

the steps of:

populating a digital repository with information corresponding to a plurality of

individual medical device components:

populating the digital repository with patient historical information associated with

a patient:

interviewing the patient having a need for a medical device to determine at least

one patient attribute;

storing the at least one patient attribute in a memory via a digital communication

link:

querying the digital repository for a subset of medical device components based

on the at least one patient attribute, the subset of medical device components

collectively forming a medical device meeting the need of the patient;

ordering the medical device over the digital communication link; and

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storing information corresponding to the medical device in the digital repository

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associated with the patient.

Claim 66 (original): The method of Claim 65, wherein the medical device comprises at

least one of a lower extremity prosthetic device, an upper extremity prosthetic device, a

lower extremity orthotic device, an upper extremity orthotic device, and a spinal orthotic

device.

Claim 67 (previously presented): The method of Claim 65, wherein the patient historical

information comprises at least one of reimbursement information and L code

information.

Claim 68 (original): The method of Claim 65, further comprising the step of:

sharing information in the digital repository with an external system.

Claim 69 (original): The method of Claim 68, wherein the external system comprises at

least one of a patient management system a billing system, and an insurance

reimbursement system.

Claim 70 (withdrawn): A system for collecting data for use in configuring a medical

device, comprising:

a digital repository populated with entries defining patient interview questions, the

entries each associated with a medical device type and having

a medical device type indicator, and

at least one patient interview question indicator;

a processor; and

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a computer readable medium encoded with processor readable instructions that

when executed by the processor implement a patient interview mechanism configured

to

provide a practitioner with access to entries in the digital repository.

allow the practitioner to provide at least one patient interview answer indicator

corresponding to a patient interview question, and

store the at least one patient interview answer in a memory.

Claim 71 (withdrawn): The system of Claim 70, wherein the medical device comprises

at least one of a lower extremity prosthetic device, an upper extremity prosthetic device,

a lower extremity orthotic device, an upper extremity orthotic device, and a spinal

orthotic device.

Claim 72 (withdrawn): The system of Claim 70, wherein at least a portion of the patient

interview mechanism is accessible via a personal data assistant.

Claim 73 (withdrawn): The system of Claim 70, wherein the digital repository comprises

a database.

Claim 74 (withdrawn): The system of Claim 70, wherein the patient interview

mechanism is further configured to accept the at least one patient interview answer

indicator from an external device.

Claim 75 (withdrawn): The system of Claim 74, wherein the external device is at least

one of a digitizer, a digital camera, and a digital video camera.

Claim 76 (withdrawn): A method for collecting data for use in configuring a medical

device, comprising the steps of:

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populating a digital repository with information corresponding to a plurality of

patient questions related to configuring a medical device:

interviewing a patient having a need for a medical device to determine at least

one patient attribute; and

storing the at least one patient attribute in a memory.

Claim 77 (withdrawn): The method of Claim 76, wherein the interviewing step comprises

inputting the at least one patient attribute via a personal data assistant.

Claim 78 (withdrawn): The method of Claim 76, wherein the interviewing step comprises

entering the at least one patient attribute via at least one of a digitizer, a digital camera,

and a digital video camera.

Claim 79 (withdrawn): The method of Claim 76, wherein the medical device comprises

at least one of a lower extremity prosthetic device, an upper extremity prosthetic device.

a lower extremity orthotic device, an upper extremity orthotic device, and a spinal

orthotic device.

Claim 80 (original): The system of Claim 3, wherein:

the customization mechanism is further configured to store a customization result

in the digital repository indicating a change made to the subset of entries selected by

the configurator mechanism, and

the computer readable medium is further encoded with processor readable

instructions that when executed by the processor further implements an algorithm

adjustment mechanism configured to adjust an algorithm of the configurator mechanism

based on the customization result stored in the digital repository, the adjustment

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causing the configurator mechanism to select a different subset of entries based on the

at least one patient interview answer indicator.

Claim 81 (original): The system of Claim 80, wherein the algorithm adjustment

mechanism comprises an application of artificial intelligence.

Claim 82 (original): The method of Claim 31, further comprising the steps of:

customizing at least one of the subset of medical device components to create a

customized medical device further meeting the need of the patient;

storing a customization result of the customizing step in the digital repository;

comparing the customization result to the subset of medical device components

to identify a customization trend; and

adjusting an algorithm used in the querying step based on the customization

trend causing a different subset of medical device components to be gueried based on

the at least one patient attribute.

Claim 83 (withdrawn): The computer program product of Claim 53, further comprising:

a fifth computer code device configured to adjust an algorithm of the third

computer code device based on a result of the fourth computer code device, the

adjustment causing the third computer code device to select a different subset of

components based on the patient specific information.

Claim 84 (withdrawn): The computer program product of Claim 83, wherein the result of

the fourth computer code device is stored in the digital repository.

Claim 85 (withdrawn): The computer program product of Claim 83, wherein the fifth

computer code device comprises an application of artificial intelligence.

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# **EVIDENCE APPENDIX**

None.

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**RELATED PROCEEDINGS APPENDIX** 

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There are no other proceedings relating to the present appeal.